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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : C07C 50/18, 50/34, A61K 31/03 A61K 31/09, 31/12, 31/135	A1	(11) International Publication Number: WO 90/08759
		(43) International Publication Date: 9 August 1990 (09.08.90)

(21) International Application Number: PCT/US90/00487

(22) International Filing Date: 26 January 1990 (26.01.90)

(30) Priority data:
302,836   27 January 1989 (27.01.89)   US

(60) Parent Application or Grant
  (63) Related by Continuation
  US   302,836 (CIP)
  Filed on   27 January 1989 (27.01.89)

(71) Applicant *(for all designated States except US)*: SLOAN-KETTERING INSTITUTE FOR CANCER RESEARCH [US/US]; 410 East 62nd Street, New York, NY 10021 (US).

(72) Inventors; and
(75) Inventors/Applicants *(for US only)* : WATANABE, Kyoichi, A. [JP/US]; 28 Wilton Road, Rye-Brook, NY 10573 (US). KOYAMA, Masao [JP/JP]; 509-3, Minowa-cho, Kouhoku-ku, Yokohama-shi, Kanagawa 223 (JP).

(74) Agent: WHITE, John, P.; Cooper & Dunham, 30 Rockefeller Plaza, New York, NY 10112 (US).

(81) Designated States: AT (European patent), BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent), US.

Published
With international search report.

(54) Title: DERIVATIVES OF CHRYSOPHANOL

(I)

(57) Abstract

The present invention concerns compounds of formula (I) wherein R^1 is hydrogen, a hydroxy group or a methoxy group; R^2 is hydrogen or a methyl group; R^3 is hydrogen or a methyl group; Y is a secondary amino group (NHalkyl) or a tertiary amino group (N(alkyl)$_2$); and Z is hydrogen or a halogen. The invention further concerns pharmaceutical compositions which comprise the above-identified compound or the acid salts thereof, and the use of the compound or compositions for treating a malignancy in a subject.

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DERIVATIVES OF CHRYSOPHANOL

The invention described herein was made in the course of
work under Grant Nos. CA-08748 and CA-18856 from the
National Cancer Institute, National Institutes of Health,
U.S. Department of Health and Human Services. The U.S.
5   Government has certain rights in this invention.

Background of the Invention

This application is a continuation-in-part of U.S. Serial
10 No. 302,836, filed January 27, 1989, the contents of which
are hereby incorporated by reference into the present
disclosure.

Some of the information set forth herein has been published
15 (see Masao Koyama, T. Ross Kelly and Kyoichi A. Watanabe,
Novel Type of Potential Anticancer Agents Derived from
Chrysophanol and Emodin.   Some Structure-Activity
Relationship Studies, [Part 1] Journal of Medicinal
Chemistry, 31:283-284 (1988) which was distributed by the
20 publisher on January 29, 1988; and Masao Koyama, Kiyobumi
Takahashi, Ting-Chao Chou, Zbigniew Darzynkiewicz, Jan
Kapuscinski, T. Ross Kelly, and Kyoichi A. Watanabe,
Intercalating Agents with Covalent Bond Forming Capability.
A Novel Type of Potential Anticancer Agents. [Part 2]
25 Derivatives of Chrysophanol and Emodin, Journal of Medicinal
Chemistry, 32:1594 (1989)).

A number of analogues of certain antitumor intercalating
agents, such as ellipticine (Le Pecq, J.-B., et al., Proc.
30 Natl. Acad. Sci. U.S.A. 71:5078 (1974); Guthrie, R. W., et
al., J. Med. Chem 18:755 (1975)) 4'-(9-acridinyl-
amino)methanesulfon-m-aniside (m-AMSA, amsacrine), (Denny,
W.A., et al., J. Med. Chem. 25:276 (1982)) and anthracycline
antibiotics (e.g., doxorubicin) (Mosh r, C.W., et al., J.
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Med. Chem. 25:18 (1982); Seshadri, R., et al., J. Med. Chem. 26:11 (1983); Myers, C. Cancer Chemother. 8:52 (1986)) have been synthesized in order to gain better therapeutic potential. However, preliminary screening data show that there is no straightforward structure-activity relationship within each group. These results seem to suggest that

5   although intercalation may be a necessary condition, it may not be sufficient and other factors may be involved that per se potentiate the anticancer activity.

Studies on the mechanism of anticancer action of antibiotic

10   CC1065 (Chidester, C.G., et al., J. Am. Chem. Soc., 103: 7629 (1981); Kanatomo, S., et al., Chem. Pharm. Bull., 29:229 (1981), Li, L.H. et al., Cancer Res. 42:999 (1982)) show that it binds to the minor groove of DNA by non-intercalative means and then slowly alkylates the amino

15   group of adenine by opening the cyclopropane ring in the antibiotic molecule. With CC1065, covalent binding of the drug with DNA, therefore, seems to be important for its potent cytotoxic activity. Mere physical interaction between the drug and DNA may not be sufficient.

20
These considerations point to the development of intercalators with slow alkylating capability. Such intercalators will bind covalently and hopefully should eventually disrupt the DNA function.

25
The compounds of the present invention have both intercalating and alkylating functionalities, and as such are potential anticancer agents.

30   The compounds of this invention may also be useful as biochemical probes for biological reactions essential for DNA synthesis.

Recent studies indicate that m-AMSA inhibits the

35   topoisom rization and catenation r actions of DNA

topoisomerase II (Wang, J.C. Annu. Rev. Biochem, 54:665 (1985)), probably by trapping the enzyme-DNA complexes. (Nelson, E.M., et al., Proc. Natl. Acad. Sci. U.S.A. 81:1361 (1984)); Chen, G.L., et al., J. Biol. Chem. 259:13560 (1984)). Other substances, such as etoposide (VP-16), adriamycin, and ellipticine (Kuhn, K.W., et al., Natl.

5   Cancer Inst. Monogr. 4:61 (1987)) also stabilize the cleavable complex between DNA topoisomerase II and DNA.

In the present invention, we show that the incorporation of an alkylating group into some DNA intercalating agents

10   greatly enhances their antileukemic properties.

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Summary of the Invention

The present invention concerns compounds of the formula:
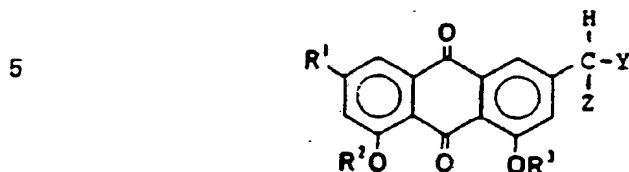
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wherein R¹ is hydrogen, a hydroxy group or a methoxy group;

R² is hydrogen or a methyl group;

R³ is hydrogen or a methyl group;

Y is a halogen, secondary amino group or a

15 tertiary amino group and

Z is hydrogen or a halogen.

The invention further concerns pharmaceutical compositions which comprise the above-identified compound or the acid

20 salts thereof, and the use of the compound or compositions for treating a malignancy in a subject.

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Brief Description of the Figures

Figure 1:   Derivatives of chrysophanol and emodin.

Figure 2:   Visible light absorption spectrum of
             derivative XIb-6 (11.3 mM in the buffer
             containing 0.01 M NaCl) alone (solid line) and
             in the presence of 0.2 mM calf thymus DNA
             (Sigma type 1) (broken line).

Figure 3:   Biological effects of chrysophanol
             derivatives.

Figure 4:   Topo II cleavable complex formation.
             (Compounds SK-31833; SK-31824; SK-31660).

Figure 5:   Topo II cleavable complex formation.
             (Compounds SK-31694; SK-31669; SK-31662).

Figure 6:   Topo II cleavable complex formation.
             (Compounds SK-31661; SK-31671; SK-31690).
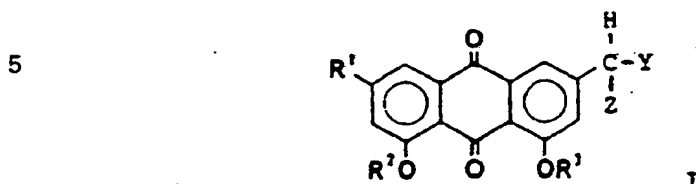
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Detailed Description of the Invention

The present invention concerns compounds of the formula (I):

$$R^1 \quad \text{anthraquinone core with} \quad \begin{array}{c} H \\ | \\ C-Y \\ | \\ Z \end{array}$$

R'O O OR'   I

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wherein R^1 is hydrogen (H), a hydroxy group (OH) or a
methoxy group (OMe);

R^2 is hydrogen or a methyl group (Me);

R^3 is hydrogen or a methyl group;

Y is a halogen, a secondary amino group or a
tertiary amino group and

Z is hydrogen or a halogen.

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When Y is a secondary amino group, e.g. [-NHalkyl], or a
tertiary amino group, e.g. [-N(alkyl)$_2$], it is preferred
that the alkyl groups be lower alkyl groups, e.g. groups
having from one to about five carbon atoms. Particularly
effective are methyl or ethyl groups. The lower alkyl
groups may also have substituents on the carbon atoms for
the hydrogens. The alkyl groups may be substituted with one
or more hydroxyl group(s), for example 2-hydroxyethyl, or
formed into organic acyl esters, such as acetyl, benzoyl or
methanesulfonyl esters, i.e. the substituents are benzoxy,
acetyloxy, or methylsulfonyloxy. Further, the alkyl groups
may be substituted with halogens(s), such as chlorine and/or
bromine to form groups such as a 2-chloroethyl or a 2-
bromoethyl.

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The invention also provides for pharmaceutical compositions
for th treatment of a malignancy in a subject comprising

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the compound or a pharmaceutically acceptable acid salt
thereof, and a pharmaceutically acceptable carrier, the
amount of the composition being an amount effective to
suppress the growth of the malignancy, preferably from 1-200
mg/kg of the body weight of the subject.

5   The invention further provides for a method of treating a
     subject have a malignancy which comprises administering to
     the subject an effective amount of the compound to suppress
     the growth of the malignancy. A subject may be any warm-
     blooded animal, preferably human.   The malignancy is
10   preferably a tumor or leukemia.

     The following Experimental Detail Section and Examples are
     set forth to aid in an understanding of the invention.
     These sections are not intended to, and should not be
15   construed to, limit in any way the invention set forth in
     the claims which follow thereafter.

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Experimental D tail

The present invention provides a novel class of compounds, anthraquinones, which possess covalent bonding capability. Such compounds may intercalate into DNA and then bind covalently to DNA, thereby exerting cytotoxic activity.

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EXPERIMENT 1:

The starting materials for the compounds of the present experiment are of the formula II:

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II
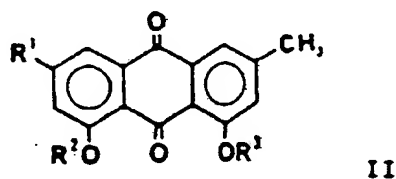
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Typical examples contain the following combinations of R^1, R^2, and R^3.

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|	R^1	R^2 and R^3	
IIa	H	H	chrysophanol
IIb	OH	H	emodin
IIc	H	Me	
IId	OMe	H	
IIe	OMe	Me	

30 Compounds IIc-IIe are known, and can be prepared readily from the natural product, chrysophanol or emodin (IIa or IIb), by the known procedures.

Compounds of formula II are treated with N-bromosuccinimide

35 (NBS) or 1,3-dibromo-5,5-dimethylhydantoin (BMH) in a

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halogenated hydrocarbon, preferably carbon tetrachloride, in the presence of a peroxid , such as m-chloroperbenzoic acid or benzoyl peroxide to give the corresponding monobromides of the formula III as the major products and dibromides of formula IV as the minor products:

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$$\text{III} \qquad \text{IV}$$
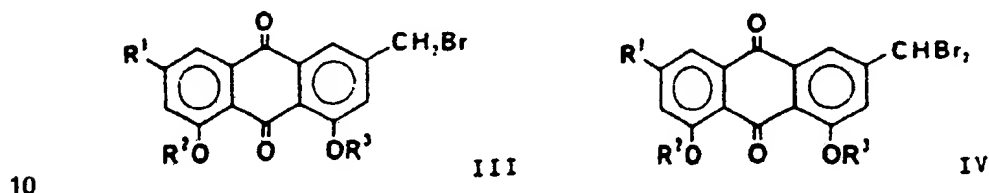
15   The reaction is carried out at a temperature range of from 25°C to 100°C, preferably at the boiling temperature of the solvent (77°C for carbon tetrachloride) in a period from one hour to three days.   The molar ratio of the reactants, formula II to NBS or BMH, can be from 1:1 to 1:3, preferably
20   1:1.2. Upon completion of the reaction, insoluble materials are removed by filtration, the filtrate concentrated, and the residue recrystallized to give formula III compounds. From the mother liquor, IV compounds can be obtained after chromatography on a silica gel column.
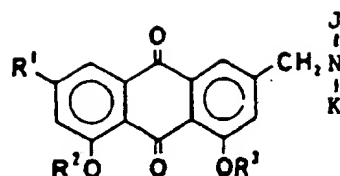25

The 1,8-dimethoxy derivatives IIIc, IIIe, IVc and IVe can be converted into the corresponding 1,8-dihydroxy derivatives IIIa, IIId, IVa, and IVd, respectively, by treatment with hydrogen bromide in acetic acid.
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Treatment of compounds of formula III with a primary or secondary amine with or without solvent affords the corresponding products of formula V:

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Some of the typical examples contain the following combinations of R^1, R^2, R^3, J and K.

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V	R^1	R^2 and R^3	J	K
Va	H	H	Et	Et
Vb	H	H	CH_2CH_2OH	CH_2CH_2OH
Vc	H	H	Et	H
Vd	H	H	CH_2CH_2OH	H
Ve	OMe	H	Et	Et
Vf	OMe	H	CH_2CH_2OH	CH_2CH_2OH
Vg	OMe	H	Et	H
Vh	OMe	H	CH_2CH_2OH	H
Vi	H	Me	Et	Et
Vj	H	Me	CH_2CH_2OH	CH_2CH_2OH
Vk	H	Me	Et	H
Vl	H	Me	CH_2CH_2OH	H
Vm	OMe	Me	Et	Et
Vn	OMe	Me	CH_2CH_2OH	CH_2CH_2OH
Vo	OMe	Me	Et	H
Vp	OMe	Me	CH_2CH_2OH	H

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The 1,8-dimethoxy derivatives of formula V (Vi-Vp) can be converted into their corresponding 1,8-

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dihydroxyanthraquinones of formula V (Va-Vh) by treatment with hydrogen bromide in acetic acid.

The 2-hydroxyethylamino derivatives (Vb, Vd, Vf, Vh, Vj, Vl, Vn and Vp) can be further converted into their corresponding 2-chloroethyl derivatives (Vq-Vx) by treatment with a conventional chlorinating agent, such as thionyl chloride, sulfonyl chloride, phosphorus oxychloride or carbon tetrachloride and triphenyl-phosphine.

V	R^1	R^2 and R^3	J	K
Vq	H	H	CH_2CH_2Cl	CH_2CH_2Cl
Vr	H	H	CH_2CH_2Cl	H
Vs	OMe	H	CH_2CH_2Cl	CH_2CH_2Cl
Vt	OMe	H	CH_2CH_2Cl	H
Vu	H	Me	CH_2CH_2Cl	CH_2CH_2Cl
Vv	H	Me	CH_2CH_2Cl	H
Vw	OMe	Me	CH_2CH_2Cl	CH_2CH_2Cl
Vx	OMe	Me	CH_2CH_2Cl	H

The reaction is carried out at a temperature range of from 0°C to 100°C, preferably at room temperature, in a period from half an hour to eight hours, in a solvent such as N,N-dimethylformamide, N,N-dimethylacetamide diethyl ether or tetrahydrofuran, preferably in N,N-dimethylformamide.

In a similar manner, the corresponding 2-bromoethyl derivatives can be obtained by bromination of the 2-hydroxyethyl intermediates with thionyl bromide, phosphorus oxybromide or carbon tetrabromide and triphenylphosphine in N,N-dimethylformamide.

Acylation of the 2-hydroxyethyl intermediates with acid anhydride, such as acetic anhydride, benzoic anhydride or

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methanesulfonic anhydride, or with acyl chloride, such as
acetyl chloride, benzoyl chloride or methanesulfonyl
chloride, in pyridine or in a mixture of chloroform and p-
dimethylaminopyridine or methylenechloride and p-
dimethylaminopyridine affords the corresponding acyl
derivatives.

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The following examples are illustrative of the process of
the products of the present experiment, but are not to be
construed as limiting.

10   EXAMPLE 1

To a hot solution of 1,8-dimethoxy-3-methyl-9,10-
anthraquinone (4.5 g, 16 mM) and 1,3-dibromo-5,5-
dimethylhydantoin (2.75 g, 19.2 mM) in carbon tetrachloride
15   (500 mL) is added benzoyl peroxide (0.7 g), and the mixture
is heated under reflux for 5 hours. The mixture is allowed
to cool to room temperature. Insoluble hydantoin is removed
by filtration, the filtrate is concentrated to dryness, and
the residue recrystallized twice from ethyl acetate to give
20   3-bromomethyl-1,8-dimethoxy-9,10-anthraquinone (3.3 g, 57%),
mp 176-178°C. ^{1}H NMR (CDCl$_3$) δ: 4.01 (3H, s, OMe), 4.03 (3H,
s, OMe), 4.52 (2H, s, CH$_2$Br), 7.26-7.84 (5H, m, aromatic H).
Analyses (C$_{17}$H$_{13}$BrO$_4$). Calculated: C, 56.53; H, 3.63; Br,
22.12. Found: C, 56.48; H, 3.67; Br, 21.93.

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The mother liquors of recrystallization of above are
concentrated, and the residue chromatographed on a silica
gel column using a mixture of benzene and ethyl acetate. 3-
Dibromomethyl-1,8-dimethoxy-9,10-anthraquinone (0.4B g) is
30   eluted form the column followed by 3-bromomethyl-1,8-
dimethoxy-9,10-anthraquinone (0.33 g). The former has the
following characteristics: mp 207-210°C, ^{1}H NMR (CDCl$_3$) δ:
4.01 (3H, s, OMe), 4.07 (3H, s, OMe), 6.67 (1H, s, CHBr$_2$),
7.26-7.92 (5H, m, aromatic H). Analyses (C$_{17}$H$_{18}$Br$_2$O$_4$).

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Calculated: C, 46.40; H, 2.75; Br, 36.31. Found: C, 46.63; H, 2.87; Br, 36.23.

Following the same procedure but using the corresponding anthraquinones as the starting materials, the following 3-bromomethyl- and 3,3-dibromomethyl-9,10-anthraquinones are
5   prepared:
3-Bromomethyl-1,6,8-trimethoxy-9,10-anthraquinone,
3-Dibromomethyl-1,6,8-trimethoxy-9,10-anthraquinone,
3-Bromomethyl-1,8-dihydroxy-9,10-anthraquinone,
3-Bromomethyl-1,8-dihydroxy-9,10-anthraquinone,
10   3-Bromomethyl-1,6,8-trihydroxy-9,10-anthraquinone,
3-Dibromomethyl-1,6,8-trihydroxy-9,10-anthraquinone,
3-Bromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone,
3-Dibromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone.

15   By the following the same procedure, but using N-bromosuccinimide   instead   of   3,3-dibromo-5,5-dimethylhydantoin, the same products above are prepared form their corresponding 3-methyl-9,10-anthraquinone starting materials.
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EXAMPLE 2

A mixture of 3-bromomethyl-1,8-dimethoxy-9,10-anthraquinone (1.05 g), 30% hydrogen bromide in acetic acid (5 mL) and
25   acetic acid (50 mL) is heated at 100°C for 5 hours. After cooling the mixture, 3-bromomethyl-1,8-dihydroxy-9,10-anthraquinone is collected by filtration, washed with acetic acid, and then air-dried to give 879 mg (91%) of the product, mp 220-222°C. ^{1}H NMR (CDCl$_3$) δ: 4.47 (2H, s,
30   CH$_2$Br), 7.27-7.91 (5H, m, aromatic H), 12.02 (1H, s, OH), 12.05 (1H, s, OH). Analyses (C$_{15}$H$_9$BrO$_4$). Calculated: C, 54.08; H, 2.71; Br, 23.99. Found: C, 54.00; H, 2.92; Br, 24.16.

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By following the same procedure but using the corresponding 1,8-dimethoxy-9,10-anthraquinones, the following compounds are prepared:

3-Dibromomethyl-1,8-dihydroxy-9,10-anthraquinone,

3-Bromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone,

5   3-Dibromomethyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone.

EXAMPLE 3

A mixture of 3-bromomethyl-1,8-dihydroxy-9,10-anthraquinone

10   (456 mg) and bis(2-hydroxyethyl)amine (600 mg) in N,N-dimethylformamide (20 mL) is stirred for 2 hours, and then partitioned between chloroform (100 mL) and water (100 mL). The organic layer is separated, washed with water (50 mL x 3), dried over sodium sulfate, and then concentrated to

15   dryness. The residue is chromatographed on a silica gel column using chloroform-methanol (15:1 v/v) as the eluent. Upon concentration of the major fraction, 3-[N,N-bis(2-hydroxyethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone is obtained as a solid: ^{1}H NMR (CDCl$_3$) δ: 2.77 (4h, t,

20   NC$\underline{H}_2$CH$_2$O), 3.78 (4H, t, NCH$_2$C$\underline{H}_2$O), 3.78 (2H, s, CH$_2$N=), 7.21–7.79 (5H, m, aromatic H), 11.95 (1H, s, OH), 12.00 (1H, s, OH).

The solid is dissolved in 1N hydrochloric acid, and the

25   solvent is removed in vacuo. The crystalline hydrochloride salt (495 mg, 91%) is triturated with methanol (5 mL), mp 204–207°C (decomposition). Analyses (C$_{19}$H$_{19}$NO$_6$·HCL). Calculated: C, 57.95; H, 5.12; N, 3.56. Found: C, 58.00; H, 5.26; N, 3.37.

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By following the same procedure but using the corresponding 3-bromomethyl-9,10-anthraquinones, the following compounds and their hydrochloric acid salts are prepared:

35   3-[N,N-Diethylamino)methyl-1,8-dihydroxy-9,10-anthraquinone,

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3-[N-Ethylamino)methyl-1,8-dihydroxy-9,10-anthraquinone,

3-[N-(2-Hydroxyethyl)amino]methyl-1,8-dihydroxy-9,10-
    anthraquinone,

3-(N,N-Diethylamino)methyl-1,8-dihydroxy-6-methoxy-
    9,10-anthraquinone,

3-[N,N-Bis(2-hydroxyethyl)amino]methyl-1,8-dihydroxy

5   -6-methoxy-9,10-anthraquinone,

3-(N-Ethylamino)methyl-1,8-dihydroxy-6-methoxy-
    9,10-anthraquinone,

3-[N-(2-Hydroxyethyl)amino]methyl-1,8-dihydroxy-6-methoxy-
    9,10-anthraquinone,

10   3-(N,N-Diethylamino)methyl-1,8-dimethoxy-9,10-
    anthraquinone,

3-[N,N-Bis(2-hydroxethyl)amino]methyl-1,8-dimethoxy-
    9,10-anthraquinone,

3-(N-Ethylamino)methyl-1,8-dimethoxy-9,10-anthraquinone,

15   3-[N-(2-Hydroxyethyl)amino]methyl-1,8-dimethoxy-9,10-
    anthraquinone,

3-(N,N-Diethylamino)methyl-1,6,8-trimethoxy-9,10-
    anthraquinone,

3-[N,N-Bis(2-hydroxethyl)amino]methyl-1,6,8-trimethoxy-

20   9,10-anthraquinone,

3-(N-Ethylamino)methyl-1,6,8-trimethoxy-9,10-anthraquinone,

3-[N-(2-Hydroxyethyl)amino]methyl-1,6,8-trimethoxy-9,10-
    anthraquinone,

25   EXAMPLE 4

To a solution of 3-[N,N-bis(2-hydroxyethyl)amino]methyl-1,8-
dihydroxy-9,10-anthraquinone,   (162   mg)   in   dry   N,N-
dimethylformamide (5 mL) is added thionyl chloride (0.2 mL).
30   After   2   hours   at   room   temperature,   the   mixture   is
concentrated   in   vacuo   to   dryness,   and   the   residue   is
triturated   well   with   methanol   (3   mL).   3-[N,N-Bis(2-
chloroethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone
that   is   crystallized   is   collected   by   filtration,   172   mg
35   (96%), mp 211-214°C (decomposition). ^{1}H NMR (CDCl$_3$) δ: 3.48

(4H, t, NC$\underline{H}_2$CH$_2$Cl), 3.85 (4H, t, NCH$_2$C$\underline{H}_2$Cl), 4.34 (2H, d, CH$_2$N=), 7.32-7.95 (5h, M, aromatic H). Analyses (C$_{19}$H$_{17}$Cl$_2$NO$_4$.HCl). Calculated: C, 52.98; H, 4.21; Cl, 24.24; N, 3.25. Found: C, 52.79; H, 4.32; Cl, 24.41; N, 3.36.

By following the same procedure but using the corresponding (2-hydroxyethyl)amino derivatives, the following compounds are prepared:

3-[N-(2-Chloroethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone,

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-6 methoxy-9,10-anthraquinone,

3-[N-(2-Chloroethyl)amino]methyl-1,8-dihydroxy-6-methoxy-9,10-anthraquinone,

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dimethoxy-9,10-anthraquinone,

3-[N-(2-Chloroethyl)amino]methyl-1,8-dimethoxy-9,10-anthraquinone,

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8,6-trimethoxy-9,10-anthraquinone.

3-[N-(2-Chloroethyl)amino]methyl-1,6,8-trimethoxy-9,10-anthraquinone,

Experimental Discussion

Table I list typical results supporting the use of the present compounds as anti-cancer agents in the treatment of subjects.

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Table I.   Inhibitory activity of some 9,10-anthraquinone
                derivatives.

Compounds	melting point (°C)	ID_{50} (μg/mL)	ID_{50} (μM)
IIa	194-195	>100	>390
IIb	256-257	>100	>335
Va (HCl)	235-238 (dec)	0.99	2.8
Vb (HCl)	204-207 (dec)	2.33	5.9
Vc (HCl)	>275	0.26	0.77
Vd (HCl)	255-261 (dec)	0.066	0.16
Ve (HBr)	240-241 (dec)	0.51	1.16
Vf (HCl)	225-227 (dec)	5.80	13.7
Vg (HBr)	>275	0.18	0.44
Vh (HCl)	259-260 (dec)	0.072	0.19
Vi (HCl)	154-158	52.3	128.3
Vj (HCl)	202-205 (dec)	>21	>49.8
Vk (HCl)	254-255 (dec)	1.30	3.59
Vl (HCl)	251-252 (dec)	53.7	142.0
Vm (HCl)	222-223 (dec)	2.90	6.91
Vn (HCl)	226-227 (dec)	12.5	27.7
Vo (HCl)	267-269 (dec)	1.44	3.67
Vp (HCl)	252-253 (dec)	5.10	12.5
Vq (HCl)	211-214 (dec)	0.058	0.13
Vr (HCl)	255-261 (dec)	2.37	7.11
Vs (HCl)	203-206 (dec)	0.010	0.023
Vu (HCl)	208-209 (dec)	5.73	12.5
Vw (HCl)	200-201 (dec)	1.30	2.66

The starting materials, chrysophanol and emodin (IIa and
IIb), are capable of intercalating into DNA but do not
possess covalent bond forming capability, and exhibit little
anticancer activity. The 1,8-dimethoxy intermediates (Vu-
Vx), that bear alkylating potential but are incapable of
intercalating into DNA due to the presence of bulky methoxy
groups, are active to a small extent against mouse leukemia
L1210 cells.   Those compounds that are capable of
intercalating into DNA and bind covalently to the DNA after
intercalation (Vq-Vt) do exhibit extremely potent activity
against L1210 cells.

Table II lists additional results supporting the anti-cancer
use of the present experiment.

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Table II.   Antileukemic   activity   of   chrysophanol
derivatives bearing alkylating potential.

J and K	R^1	R^2	R^3	L1210: $ID_{50}, \mu M$
Et	H	Me	Me	128.3
Et	H	H(or Me)	Me (or H)	11.9
Et	H	H	H	2.75
CH_2CH_2OH	H	Me	Me	>50
CH_2CH_2OH	H	H(or Me)	Me (or H)	23.4
CH_2CH_2OH	H	H	H	5.92
CH_2CH_2Cl	H	Me	Me	12.5
CH_2CH_2Cl	H	H(or Me)	Me (or H)	1.40
CH_2CH_2Cl	H	H	H	0.13
Et	OMe	Me	Me	6.91
Et	OMe	H	H	1.16
CH_2CH_2OH	OMe	Me	Me	>27.6
CH_2CH_2OH	OMe	H	H	13.7
CH_2CH_2Cl	OMe	Me	Me	2.66
CH_2CH_2Cl	OMe	H(or Me)	Me (or H)	1.75
CH_2CH_2Cl	OMe	H	H	0.023

3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-9,10-
anthraquinone (Vq) is extremely potent against mouse
leukemia L1210 made resistant to Cisplatin. At the dosage
of 100 mg/kg/day x 5 (ip), mice inoculated with Cisplatin
resistant leukemia L1210/Cisplatin are cured.

The process of treating tumors according to this invention
comprises administering to a subject having an abnormal
proportion f leukocytes or other evidence of a malignancy,
a therapeutic nontoxic amount of a compound of the

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experiment such as 3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-9,10-anthraquinone, as such or in the form of a pharmaceutically acceptable salt thereof. The invention also provides a pharmaceutical composition in dosage unit form comprising from 1 to 200 mg/kg of a compound of the invention, per dosage unit, together with pharmaceutically

5   acceptable nontoxic inert carrier or diluent thereof as described above. A subject may be any warm-blooded animal and is preferably human.

EXPERIMENT 2:

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Fifty-one new $\underline{C}$-methyl-modified derivatives of the anthraquinones chrysophanol and emodin or their various methyl ethers were prepared for structure-activity relationship studies of anticancer activity against mouse

15   leukemia L1210 and human leukemia HL-60 cells. Representative compounds were spectrophotometrically studied for their capacity to interact with natural and denatured DNA. In general, those anthraquinones bearing an amino function interact with DNA. 1,8-Dimethoxyanthraquinones are

20   incapable of intercalating into DNA. 1- or 8-Monohydroxy-monomethoxyanthraquinones, however, interact with DNA to some extent. No straightforward correlation is apparent between the DNA-affinity data of the compounds studied spectrophotometrically and their cytotoxic effects.

25   Cytotoxic potencies of these compounds on cell growth inhibition during a 72-h period are inversely correlated to their potencies when inhibiting [^{3}H]TdR incorporation into DNA during the initial 30 min of exposure. Surprisingly, some compounds that showed more cytotoxicity did not inhibit

30   initial TdR incorporation (0-30 min), while some others that strongly inhibited TdR incorporation initially did not exhibit cytotoxicity in 72 h. The results suggest that the cytotoxicity produced by these compounds is time dependent and is not a direct result of initial inhibition of DNA

35   replication.

It is well-known that one the of the metabolites of ellipticine, 9-hydroxyellipticine (Lesca, P., et al., Biochem. Pharmacol. 26:2169 (1977)) (9-OH-E), is also a potent anticancer agent. (Le Pecq, J.-B.; Cancer Res. 36:3067 (1976)) 2-N-Methyl-9-hydroxyellipticinium (9-OH-NME) is one of the most active drugs among the ellipticine

5 analogues (Bernadou, J. Proc. Natl. Acad. Sci. U.S.A. 81:1297 (1984)). The latter is easily oxidized by peroxidases to 9-oxo-2-methylellipticinium (Auclair, C., et al., J. Med. Chem. 24:289 (1981); Bernadou, J., et al., J. Med. Chem. 26:574 (1983)) (9-oxo-NME), which is highly

10 electrophilic and alkylates various nitrogen, (Meunier, G.; Tetrahedron Lett. 26:574 (1983); Auclair, C.; J. Med. Chem. 27:1161 (1984)) sulfur, (Monsarrat, B.; Biochem. Pharmacol. 32:3887 (1983)) and oxygen (Bernadou, J., et al., J. Med. Chem. 26:574 (1983); Meunier, G., et al., Tetrahedron Lett.

15 26:574 (1983)) nucleophiles. Among biological macromolecules, proteins, (Auclair, C., et al., Biochem. Pharmacol. 32:3883 (1984) polyadenylate, (Dugue, B., et al., Biochem. Biophys. Res. Commun. 124:416 (1984)) RNA, (Dugue, B., et al.) and DNA (Auclair, C., et al., Biochemistry

20 25:1240 (1986) are easily alkylated by 9-oxo-NME. This "biooxidative alkylation" has been proposed as a possible mode of anticancer action. (Dugue, B., et al., Cancer Res. 46:3828 (1986).

25 For other intercalating anticancer agents, such as amsacrine (m-AMSA), and the anthracycline antibiotics, extensive studies on their mechanism of anticancer action (Nelson, E.M., et al., Proc. Natl. Acad. Sci. U.S.A. 81:1361 (1984); Chen, G.L., et al., J. Biol. Chem. 259:13560 (1984); Riou,

30 J.F., et al., Biochem. Pharmacol. 35:4409 (1986); Crooke, S.T., Reich, S.D.; Eds Anthracyclines: Current Status and New Developments; Academic Press: New York, (1980); Myers, C.E. In Pharmacologic Principles of Cancer Treatment; Chabner, B., Ed.; Saunders: Philadelphia p. 416 (1982);

35 DiMarco, A. In Cancer Medicine; Holland, J.F., Frei, E. Eds;

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Lea and Febiger: Philadelphia, 2nd ed., p. 872 (1982);
Muggia, F.M., Young, C.W., Carter, S.K. Eds. Anthracycline
Antibiotics in Cancer Therapy; Martinus Nijhoff:
Hague/Boston/New York, pp. 71-174 (1982)) and QSAR studies
directed toward the development of more selective drugs have
been conducted. (Ferguson, L.R., Denny, W.A., J. Med. Chem.
5   23:269 (1980); Denny, W.A., et al., J. Med. Chem. 25:276
(1982); Alexander, J., et al., J. Med. Chem. 27:1343
(1984)). Whether these intercalators bind covalently to
biomolecules has not been established.

10   In order to test our hypothesis that intercalating agents
with covalent bond forming capability may exert potent
cytocidal activity, we chose chrysophanol Ia and emodin Ib
(Figure 1) as the starting materials; Ia and Ib were
isolated from crude rhubarb extract. (Kelly, T.R.; J. Org.
15   Chem. 48:3573 (1983)). Both compounds exhibit little
anticancer or cytocidal activity. Their structural aromatic
features indicate that these compounds and their derivatives
(particularly positively charged ones) may intercalate into
double helix of nucleic acids.
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Chemistry

The hydroxy groups at the 1- and 8-positions of I were
methylated with methyl sulfate and K_2CO_3 in acetone
25   (Alexander, J., et al., J. Org. Chem. 45:20 (1980)) to the
known 1,8-dimethoxy-9,10-anthraquinones II. The $\underline{c}$-methyl
group of II was then brominated with NBS (Banvill, J., et
al., J. Chem Soc. Perkin Trans. I, 613 (1976)) or 1,3-
dibromo-5,5-dimethylhydantoin (BDH) (Sargent, M.V., et al.,
30   J. Chem. Soc., C 2763 (1969); Cava, M.P., et al.,
Tetrahedron 40:4767 (1984)) in carbon tetrachloride in the
presence of benzoyl peroxide to give monobromide III as the
major product along with a samll amount of dibromide IV.
Treatment of III with various amines including mono(2-
35   hydroxyethyl)amine and bis(2-hydroxy thyl)amine afforded the

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corresponding alkyl-amino derivatives IX (Table III).
Chlorination of the (2-hydroxyethyl)amino derivatives IX-2
gave the corresponding 3-[[(2-chloroethyl)amino]methyl]-
9,10-anthraquinones IX-3.   On the basis of reports by
Anderson et al. (Anderson, W.K., et al., J. Med. Chem.
20:812 (1977); Anderson, W.K., et al., J. Med. Chem. **22**:977

5   (1979); Anderson, W.K., et al., J. Med. Chem. **26**:1333
(1983); Zwelling, L.A. Cancer Metastasis Rev. **4**:263 (1985))
that certain carbamates are susceptible to nucleophilic
attack, we synthesized N-methylcarbamate IX-4 by treatment
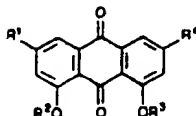of IX-2 with N-methyl isocyanate.

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Table III

Synthetic Derivatives of Chrysophanol and Emodin

$$R^1, R^4, R^2O, OR^3$$ (anthraquinone structure)

compd[a]	R^1	R^2	R^3	R^4	mp, °C[b]	formula
IIa	H	Me	Me	Me	189–190	$C_{17}H_{14}O_4$
IIb	OMe	Me	Me	Me	228–229	$C_{18}H_{16}O_5$
IIIa	H	Me	Me	CH_2Br	176–178	$C_{17}H_{13}BrO_4$
IIIb	OMe	Me	Me	CH_2Br	250–254	$C_{18}H_{15}BrO_5$
IVa	H	Me	Me	$CHBr_2$	207–210	$C_{17}H_{12}Br_2O_4$
IVb	OMe	Me	Me	$CHBr_2$	254–257	$C_{18}H_{14}Br_2O_5$
Va	H	H or Me	Me or H	CH_2Br	213–215	$C_{16}H_{11}BrO_4$
Vb	OMe	H or Me	Me or H	CH_2Br	199–201	$C_{17}H_{13}BrO_5$
VIa	H	H	H	CH_2Br	220–222	$C_{15}H_9BrO_4$
VIb	OMe	H	H	CH_2Br	249–250	$C_{16}H_{11}BrO_5$
VIIa	H	H or Me	Me or H	$CHBr_2$	176–178	$C_{16}H_{10}Br_2O_4$
VIIb	OMe	H or Me	Me or H	$CHBr_2$	202–203	$C_{17}H_{12}Br_2O_5$
VIIIa	H	H	H	$CHBr_2$	211–213	$C_{15}H_8Br_2O_4$
VIIIb	OMe	H	H	$CHBr_2$	237–238	$C_{16}H_{10}Br_2O_5$
IXa-1	H	Me	Me	CH_2NEt_2	154–158	$C_{21}H_{25}NO_4\cdot HCl\cdot H_2O$
IXa-2	H	Me	Me	$CH_2N(CH_2CH_2OH)_2$	202–206d	$C_{21}H_{25}NO_6\cdot HCl$
IXa-3	H	Me	Me	$CH_2N(CH_2CH_2Cl)_2$	205–206d	$C_{21}H_{23}Cl_2NO_4\cdot HCl$
IXa-4	H	Me	Me	$CH_2N(CH_2CH_2OCONHMe)_2$	120d	$C_{25}H_{28}N_3O_8\cdot HCl$
IXa-5	H	Me	Me	CH_2NHEt	254–255d	$C_{19}H_{19}NO_4\cdot HCl\cdot H_2O$
IXa-6	H	Me	Me	$CH_2NHCH_2CH_2OH$	251–252d	$C_{19}H_{19}NO_5\cdot HCl$
IXa-7	H	Me	Me	$CH_2NHCH_2CH_2Cl$	208–209d	$C_{19}H_{18}ClNO_4$
IXb-1	OMe	Me	Me	CH_2NEt_2	222–223d	$C_{22}H_{27}NO_5\cdot HCl$
IXb-2	OMe	Me	Me	$CH_2N(CH_2CH_2OH)_2$	225–227d	$C_{22}H_{25}NO_7\cdot HCl$
IXb-3	OMe	Me	Me	$CH_2N(CH_2CH_2Cl)_2$	200–201d	$C_{22}H_{23}Cl_2NO_5\cdot HCl$
IXb-5	OMe	Me	Me	CH_2NHEt	267–269d	$C_{20}H_{21}NO_5\cdot HCl$
IXb-6	OMe	Me	Me	$CH_2NHCH_2CH_2OH$	252–253d	$C_{20}H_{21}NO_6\cdot HCl$
IXb-7	OMe	Me	Me	$CH_2NHCH_2CH_2Cl\cdot HCl\cdot DMF$	204–205d	$C_{20}H_{20}ClNO_5\cdot HCl\cdot C_3H_7NO$
Xa-1	H	H or Me	Me or H	CH_2NEt_2	225–227d	$C_{20}H_{23}NO_4\cdot HBr$
Xa-2	H	H or Me	Me or H	$CH_2N(CH_2CH_2OH)_2$	209–216d	$C_{20}H_{23}NO_6\cdot HCl$
Xa-3	H	H or Me	Me or H	$CH_2N(CH_2CH_2Cl)_2$	203–205d	$C_{20}H_{21}Cl_2NO_4\cdot HCl$
Xa-4	H	H or Me	Me or H	$CH_2N(CH_2CH_2OCONHMe)_2$	178–182d	$C_{24}H_{27}N_3O_8\cdot HCl$
Xb-1	OMe	H or Me	Me or H	CH_2NEt_2	110–112	$C_{21}H_{25}NO_5$
Xb-2	OMe	H or Me	Me or H	$CH_2N(CH_2CH_2OH)_2$	221–223	$C_{21}H_{25}NO_7\cdot HCl$
Xb-3	OMe	H or Me	Me or H	$CH_2N(CH_2CH_2Cl)_2$	152–154	$C_{21}H_{21}Cl_2NO_5$
XIa-1	H	H	H	CH_2NEt_2	235–238d	$C_{18}H_{18}NO_4\cdot HCl\cdot^1/_2H_2O$
XIa-2	H	H	H	$CH_2N(CH_2CH_2OH)_2$	204–207d	$C_{18}H_{18}NO_6\cdot HCl$
XIa-3	H	H	H	$CH_2N(CH_2CH_2Cl)_2$	211–214d	$C_{18}H_{17}Cl_2NO_4\cdot HCl$
XIa-4	H	H	H	$CH_2N(CH_2CH_2OCONHMe)_2$	125–131	$C_{22}H_{23}N_3O_8\cdot HCl$
XIa-5	H	H	H	CH_2NHEt	>275	$C_{17}H_{15}NO_4\cdot HCl$
XIa-6	H	H	H	$CH_2NHCH_2CH_2OH$	255–261d	$C_{17}H_{15}NO_5\cdot HCl$
XIa-7	H	H	H	$CH_2NHCH_2CH_2Cl$	255–261d	$C_{17}H_{14}ClNO_4\cdot HCl$
XIa-8	H	H	H	CH_2NH_2	240–245d	$C_{15}H_{11}NO_4\cdot HCl\cdot H_2O$
XIa-9	H	H	H	CH_2NMe_2	282–283d	$C_{17}H_{15}NO_4\cdot HCl$
XIa-10	H	H	H	$CH_2NH(CH_2)_2Me$	>275	$C_{18}H_{17}NO_4\cdot HCl$
XIa-11	H	H	H	$CH_2NHCH_2CH_2CH_2OH$	251–252d	$C_{18}H_{17}NO_5\cdot HCl$
XIa-12	H	H	H	$CH_2N(CH_2CH_2OCONH\text{-}iPr)_2$	159–160	$C_{27}H_{29}N_3O_8\cdot HCl$
XIa-13	H	H	H	$CH_2N(CH_3)_3$	255–257d	$C_{18}H_{17}NO_4\cdot HCl$
XIa-14	H	H	H	$CH_2N(CH_3)_2$	246–247d	$C_{18}H_{16}NO_4\cdot HCl$
XIa-15	H	H	H	$CH_2(imidazol\text{-}1\text{-}yl)$	270–272d	$C_{18}H_{12}N_2O_4\cdot HCl\cdot H_2O$
XIb-1	OMe	H	H	CH_2NEt_2	240–241d	$C_{20}H_{21}NO_5\cdot HBr$
XIb-2	OMe	H	H	$CH_2N(CH_2CH_2OH)_2$	225–227d	$C_{20}H_{21}NO_7\cdot HCl$
XIb-3	OMe	H	H	$CH_2N(CH_2CH_2Cl)_2$	203–206d	$C_{20}H_{19}Cl_2NO_5\cdot HCl$
XIb-5	OMe	H	H	CH_2NHEt	>275	$C_{18}H_{17}NO_5\cdot HBr$
XIb-6	OMe	H	H	$CH_2NHCH_2CH_2OH$	259–260d	$C_{18}H_{17}NO_6\cdot HCl$
XIb-7	OMe	H	H	$CH_2NHCH_2CH_2Cl$	powder	$C_{18}H_{16}ClNO_5$

[a] All the compounds were analyzed for C, H, X (Br or Cl), and/or N. Analyses for these elements were within ±0.4% of the theoretical values required unless specified otherwise. [b] For nitrogen-containing compounds, melting points were of the HX salt. [c] d = decomposition. [d] Unstable, and satisfactory analyses could not be obtained.

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The methyl protecting groups at 1 and 8 could be removed
stepwise at various stages (Figure 1). Thus, treatment of
IX with HBr in acetic acid at room temperature afforded a
crystalline mixture of 1-O-methyl- and 8-O-methyl-
anthraquinones X, whereas acid hydrolysis at reflux

5   temperature for a few hours resulted in complete
demethylation, giving rise to XI. Later, it was found that
partially methylated chrysophanol and emodin could be
directly brominated to the corresponding mixtures of the
monobromides V (major products) and dibromides VII. The

10   former were treated with amines to give X, which were
further converted to the corresponding XI. Alternatively,
the 3-[(alkylamino)methyl] derivatives XI were prepared by
amination of VI. Chlorination of XIa-2 with thionyl
chloride afforded XIa-3. N-Methyl- and N-isopropyl-

15   carbamates, XIa-4 and XIa-12, respectively, were prepared by
treatment of XIa-2 with the corresponding N-alkyl
isocyanates.

Spectrophotometric Studies for DNA Interactions

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Some representative compounds were studied for their ability
to interact with nucleic acids in solution by comparison of
the electronic spectrum of drug alone with that of the drug
in the presence of an excess of nucleic acid. All drugs

25   studied have an absorption band in the visible region,
separate from the absorption band of the nucleic acids, and
therefore any changes in band intensity and position were
indicative of drug chromophore-DNA interaction. It was
observed that both chrysophanol (Ia) and emodin (Ib) and

30   their derivatives lacking the basic center do not appear to
interact with DNA to any significant extent. Since these
compounds have low solubility in aqueous solutions, the
sp ctral measurement alon may not be sufficient howev r to
allow one to draw a definite conclusion.

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Generally, those anthraquinones bearing an amino function (e.g., Xa-1, Xb-1, XIa-1, X1a-2, XIb-2, and XIb-6) interact with both native and thermally denatured DNA but more strongly with native DNA (Figure 2). As expected, 1-8-di-$\underline{O}$-methylchrysophanol and 1,6,8-tri-$\underline{O}$-methylemodin analogues

5   IXa and IXb do not interact with DNA. Anthraquinones wherein one peri-hydroxyl group is methylated (e.g., Xa-2 and Xb-2) interact with DNA to a lesser extent than the correspnding unmethylated (XIa-2 and XIb-2). It is interesting to note that while DNA induced changes in absorption spectra of some derivatives (e.g., XIa-7 and XIb-

10   7), these do not appear to connected with the alklating capabilities in view of the fact that they could be reversed by addition of Me_2SO (to 1:1 v/v). The dissociation of nonbonded ligand-DNA complexes in the presence of organic solvents is a phenomenon well documented. (Wakelin, L.P.G.,

15   et al., Biochemistry 20:5779 (1981)).

Changes in the absorption spectra of the drugs (Figure 2) are not inconsistent with the possiblity of the intercalative mode of binding. Other types of nonbinding

20   interactions, however, (e.g. binding to the minor groove of the double helix (Bloomfield, V.A., et al., In Physical Chemistry of Nucleic Acids; Harper and Row: New York, pp. 432-434 (1974); Jorgenson, K.F., et al., J. Biomol. Struct. Dyn. 6:1005 (1988)) cannot be excluded. It is well-known

25   that intercalative binding, which most often has an ionic component, is affected by a rise in concentration of salts, (Kapuscinski, J., Darzynkiewicz, Z.J., Biomol. Struct. Dyn. 5:127 (1987), and compounds such as Xa-2, Xb-2, XIb-3, and XIb-7 lost the ability to interact with DNA when Na^+

30   concentration was increased from 0.01 to 0.1 M. On the basis of this fact one can conclude that the affinity for DNA of these compounds is lower than those that interact with DNA at both ionic strengths.

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No straightforward correlation is apparent between the DNA-affinity data of the drugs studied and their biological activity.

Biological Activities

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Preliminary biological data for inhibiting cell growth of murine L1210 leukemic cells and human acute promyelocytic leukemia cells (HL-60) during 72 h of exposure to the compounds are given in Table IV. The potencies for inhibiting [³H]TdR incorporation into DNA in HL-60 cells

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during the initial 30-min period are also given in Table IV. It is interesting to note that 1,8-di-$\underline{o}$-methyl derivatives are uniformly devoid of activity against L1210 leukemic cells. These results are consistent with published data that compare the ratios of the potencies (ID_{50}'s) for cell growth inhibition (A) and inhibition of [³H]TdR

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incorporation into DNA in HL-60 cells (B). The B/A ratios allow an indirect estimation of whether or not cytotoxicities exerted by these analogues are primarily due to initial inhibition of DNA synthesis. The B/A ratios for Vb, VIIa, VIIb, VIIIb, Xa-3, and XIa-3 are 41, 217, 237,

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579, 39, and 383, respectively, suggesting that these compounds exert their initial effects mainly on processes other than DNA synthesis per se (Table V). These results suggest that these compounds exert their cytotoxic effects in a time-dependent manner and their initial action is

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targeted at the sites other than DNA elongation. Whether these analogues, like $\underline{m}$-AMSA, ellipticine, or anthracyclines, act by inhibiting DNA topoisomerase II remains to be explored. It is of interest to note that the above-mentioned anthraquinones are among the most potent

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antileukemic analogues listed in Table IV, with ID_{50} values ranging from 1.4×10^{-6} to 4.2×10^{-11} for L1210[48] cells and 1.7×10^{-6} to 6.0×10^{-8} M for HL-60 cells. Our preliminary experiments indicate that the compounds arrest cells in the S and/or G_2 phases of the cell cycle (unpublished results).

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TABLE IV
Biological Activities of Derivatives of Chrysophanol and Emodin

compd	ID$_{50}$ M (L1210 cell growth)	ID$_{50}$ M (HL-60 cell growth (72 b)) (A)	ID$_{50}$ M (HL-60 TdR into DNA) (B)	B/A
IIa	2.8×10^{-4}	1.0×10^{-4}	1.9×10^{-4}	1.9
IIb	1.0×10^{-4}	4.9×10^{-4}	2.1×10^{-4}	0.43
IIIa	9.2×10^{-4}	4.4×10^{-4}	1.1×10^{-4}	2.5
IIIb	6.8×10^{-7}	8.4×10^{-4}	1.2×10^{-4}	0.14
IVa	8.9×10^{-7}	2.1×10^{-4}	9.0×10^{-4}	4.3
IVb	4.1×10^{-4}	6.0×10^{-4}	5.8×10^{-4}	9.7
Va	8.8×10^{-4}	7.1×10^{-4}	1.2×10^{-4}	1.7
Vb	6.8×10^{-4}	1.7×10^{-4}	7.0×10^{-4}	41
VIa	8.6×10^{-4}	7.1×10^{-4}	1.6×10^{-4}	2.3
VIb	5.9×10^{-4}	2.5×10^{-4}	3.4×10^{-4}	13.6
VIIa	4.2×10^{-11}	6.0×10^{-4}	1.3×10^{-4}	217
VIIb	1.0×10^{-4}	9.7×10^{-4}	2.3×10^{-4}	237
VIIIa	4.4×10^{-7}	2.5×10^{-4}	3.9×10^{-4}	15.6
VIIIb	1.0×10^{-4}	1.9×10^{-7}	1.1×10^{-4}	579
IXa-1	1.3×10^{-4}	3.0×10^{-4}	1.9×10^{-4}	6.3
IXa-2	$>5.0 \times 10^{-4}$	4.9×10^{-7}	1.4×10^{-4}	0.03
IXa-3	1.3×10^{-4}	1.8×10^{-4}	1.4×10^{-4}	7.8
IXa-4	7.2×10^{-4}	9.5×10^{-4}	9.4×10^{-4}	0.99
IXa-5	1.2×10^{-4}	4.8×10^{-4}	2.9×10^{-4}	0.60
IXa-6	1.4×10^{-4}	5.5×10^{-4}	4.6×10^{-4}	0.84
IXa-7	2.6×10^{-4}	8.6×10^{-4}	1.9×10^{-4}	0.22
IXb-1	6.9×10^{-4}	7.9×10^{-4}	4.2×10^{-4}	0.53
IXb-2	$>2.7 \times 10^{-4}$	1.0×10^{-4}	1.7×10^{-4}	0.17
IXb-3	2.7×10^{-4}	2.0×10^{-4}	8.9×10^{-4}	4.5
IXb-5	9.8×10^{-4}	1.7×10^{-4}	1.3×10^{-4}	0.76
IXb-6	1.7×10^{-4}	1.1×10^{-4}	2.6×10^{-4}	0.24
IXb-7	3.2×10^{-4}	5.8×10^{-4}	1.8×10^{-4}	3.1
Xa-1	1.2×10^{-4}	5.2×10^{-4}	1.4×10^{-4}	2.7
Xa-2	$>2.4 \times 10^{-4}$	2.1×10^{-4}	8.9×10^{-4}	4.2
Xa-3	1.4×10^{-4}	3.9×10^{-7}	1.5×10^{-4}	38.5
Xa-4	8.9×10^{-4}	1.2×10^{-4}	1.2×10^{-4}	1
Xb-1	6.9×10^{-4}	6.7×10^{-4}	1.8×10^{-4}	26.9
Xb-2	8.8×10^{-4}	$>5.0 \times 10^{-4}$	9.6×10^{-4}	0.02
Xb-3	3.3×10^{-4}	5.2×10^{-7}	5.4×10^{-4}	10.4
XIa-1	2.8×10^{-4}	1.8×10^{-4}	1.4×10^{-4}	7.8
XIa-2	5.9×10^{-4}	3.3×10^{-4}	1.4×10^{-4}	4.2
XIa-3	1.3×10^{-7}	1.8×10^{-7}	6.9×10^{-4}	383
XIa-4	$>1.8 \times 10^{-4}$	2.1×10^{-4}	2.8×10^{-4}	0.13
XIa-5	7.7×10^{-7}	8.7×10^{-4}	1.8×10^{-4}	2.1
XIa-6	1.6×10^{-7}	8.6×10^{-7}	1.5×10^{-4}	17.4
XIa-7	7.1×10^{-4}	7.5×10^{-4}	3.9×10^{-4}	5.2
XIa-8	8.5×10^{-4}	1.1×10^{-4}	2.5×10^{-4}	2.3
XIa-9	2.2×10^{-4}	2.7×10^{-4}	9.3×10^{-4}	3.3
XIa-10	4.6×10^{-4}	4.1×10^{-4}	2.5×10^{-4}	6.1
XIa-11	6.7×10^{-7}	3.2×10^{-4}	1.6×10^{-4}	5.0
XIa-12	2.7×10^{-4}	2.7×10^{-4}	1.7×10^{-4}	0.63
XIa-13	1.6×10^{-4}	2.1×10^{-4}	1.0×10^{-4}	4.8
XIa-14	4.0×10^{-4}	2.8×10^{-4}	6.3×10^{-4}	22.5
XIa-15	7.4×10^{-4}	4.8×10^{-4}	4.3×10^{-4}	9.0
XIb-1	1.2×10^{-4}	3.4×10^{-4}	3.5×10^{-4}	10.3
XIb-2	1.4×10^{-4}	6.6×10^{-4}	4.5×10^{-4}	5.2
XIb-3	2.3×10^{-4}	6.1×10^{-7}	1.2×10^{-4}	19.7
XIb-5	7.2×10^{-7}	2.1×10^{-4}	1.3×10^{-4}	6.2
XIb-6	5.0×10^{-7}	1.6×10^{-4}	1.2×10^{-4}	7.5
XIb-7	1.8×10^{-4}	9.2×10^{-4}	2.2×10^{-4}	23.9

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TABLE V

Inverse Relationship Betwe n Cell Growth Inhibition
and Inhibition of Initial Thymidine Incorporation Into
DNA in HL-60 Cells by Chrysophanol Derivatives[a]

no. of compds examined	value of IC$_{50}$ (cell growth), μM		ratio of IC$_{50}$ (dThd incorpn)/ IC$_{50}$ (cell growth),
	range	mean ± SE	mean ± SE
8	<1	0.36 ± 0.10	167.3 ± 72.4
17	1–5	2.73 ± 0.24	9.55 ± 2.30
15	5–10	7.42 ± 0.53	6.32 ± 2.10
7	10–50	28.43 ± 5.51	3.03 ± 1.83
8	>50	204.4 ± 65.4	0.22 ± 0.09

[a] Cell growth inhibition was measured at the end of 72-h exposure to each compound as described under Experimental Section. Inhibition of [^{3}H]dThd incorporation into DNA was measured during the first 30 min of exposure to each corresponding compound as described under Experimental Section.

Materials and Methods

Melting points were determined on a Thomas-Hoover capillary apparatus and are uncorrected. Elemental analyses were performed by M-H-W Laboratories, Phonenix, AZ, and all new compounds with the exception of IXa-7 and XIa-7, which were unstable, analyzed correctly. ^{1}H NMR spectra were recorded on a JEOL FX90Q spectrometer with Me$_4$Si as the internal standard. Chrysophanol (Ia) and emodin (Ib) were isolated form rhubarb extract by the procedure of Kelly et al. (Kelly, T.R., et al. J. Org. Chem. 48:3573 (1983)) except CH$_2$Cl$_2$ was used instead of Et$_2$O throughout the isolation process.

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The following examples are illustrative of the process and products of the present invention, but are not to be construed as limiting.

<u>EXAMPLE 5</u>

I,8-Dimethoxy-3-methyl-9,10-anthraquinone (1,8-Di-<u>O</u>-methylchrysophanol, IIa). A mixture of chrysophanol (Ia, 7.0 g, 0.029 mol), K_2CO_3 (10 g, 0.071 mol), and Me_2SO_4 (10 mL, 0.1 mol) in Me_2CO (300 mL) was stirred under reflux for 16 h and then concentrated in vacuo. The residue was triturated well with water (300 mL), and the crystalline IIa (7.5 g, 96%) was collected by filtration and air-dried: mp 191-193 °C (lit, (Beilstein, 8:473) mp 195 °C); ^{1}H NMR ($CDCl_3$) δ 2.46 (3 H, s, 3-Me), 3.98 (3 H, s, OMe), 3.99 (3 H, s, OMe), 7.08-7.86 (5 H, m, H-2,4,5,6,7). Anal. ($C_{17}H_{14}O_4$) C, H.

<u>EXAMPLE 6</u>

1,3,8-Trimethoxy-6-methyl-9,10-anthraquinone (1,3,8-Tri-<u>O</u>-methylemodin, IIb). In a similar manner, emodin (Ib, 1.0 g, 3.9 mmol) was converted into IIb (1.04 g, 90%): mp 225 °C (lit. (Beilstein, 8:523) mp 225 °C); ^{1}H NMR (Me_2SO-$\underline{d}_6$) δ 2.43 (3 H, s, 6-Me) 3.88 (3 H, s, OME), 3.92 (6 H, s, 2 x OMe), 6.94 (1 H, d, H-7, $\underline{J}_{5,7}$ = 2.2 Hz), 7.13 (1 H, d, H-5, $\underline{J}_{5,7}$ = 2.2 Hz), 7.33 (1 H, s, H-2), 7.46 (1 H, s, H-4). Anal. ($C_{18}H_{16}BrO_5$) C, H.

<u>EXAMPLE 7</u>

3-(Bromomethyl)-1,8-dimethoxy-9,10-anthraquinone (IIIa). To a hot solution of IIa (4.5 g, 0.016 mol) and BDH (2.75 g, 0.019 mol) in CCl_4 (500 mL) was added Bz_2O_2 (0.7 g), and the mixture was heated under reflux for 5 h. The mixture was allowed to cool room temperature. Insoluble hydantoin was removed by filtration, the filtrate was concentrated in vacuo, and the residue was crystallized twice from EtOA c to give IIIa (3.3 g, 57%): mp 176-178 °C (lit. (Blankespoor, R.L., et al., J. Org. Chem. 52:2059 (1987) mp 174-175 °C); ^{1}H NMR ($CDCl_3$) δ 4.01 (3 H, s, OMe), 4.03 (3 H, s, OME), 4.01

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(3 H, s, OMe), 4.52 (2 H, s, Ch$_2$Br), 7.26-7.84 (5 H, m, H-2,4,5,6,7). Anal. (C$_{17}$H$_{13}$BrO$_4$) C,H, Br.

EXAMPLE 8

3-(Dibromomethyl)-1,8-dimethyoxy-9,10-anthraquinone (IVa). The mother liquors of recrystallization were concentrated, and the residue was chromatographed on a silica gel column
5   using a mixture of C$_6$H$_6$ and EtOAc (3:1) to give IVa (0.48 g): mp 207-210 degrees C; ^{1}H NMR (CDCl$_3$) lambda 4.01 (3 H, s, OMe), 4.07 (3 H, s, OMe), 6.67 (1 H, s, CHBr$_2$), 7.26 (5 H, m, H-2,4,5,6,7). Anal (C$_{17}$H$_{13}$Br$_2$O$_4$) C, H, Br.

10   A further amount of IIIa (0.33 g) was eluted from the column, making the total yield of 62.7%.

EXAMPLE 9

6-(Dibromomethyl)-1,3,8-trimethoxy-9,10-anthraquinone (IVb).
15   In a similar manner, IIb (3.12 g, 0.01 mol) was brominated to give IIIb (3.06 g, 74.4%) [mp 250-254 °C; (lit. Alexander, J., et al., J. Org. Chem. 45:20 (1980)) mp. 233.5-234 degrees C); ^{1}H NMR (CDCl$_3$) was identical with that reported (Alexander, J., et al.)] and IVb (197 mg) [mp 254-
20   257 degrees C; ^{1}H NMR (CdCl$_3$) lambda 3.95 (6 H, s, 2 X OMe), 3.97 (3 H, s, OMe), 6.67 (1 H, s, CHBr$_2$), 6.78 (1 H, d, H-7, $J_{5,7}$ = 2.47 Hz), 7.32 (1 H, d, H-5), 7.54 (1 H, d, H-2, $J_{2,4}$ = 1.92 Hz), 7.90 (1 H, d, H-4)]. Anal. (C$_{17}$H$_{14}$Br$_2$O$_5$) C, H, Br.

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EXAMPLE 10

3-(Bromomethyl)-1(and 8)-Hydroxy-8(and 1)-Methoxy-9,10-anthraquinone (Va). A mixture of IIIa (70 mg, 0.19 mmol in HOAc (10 mL) and 30% HBr/HOAc (1 mL) was stirred overnight at room temperature and then concentrated in vacuo. The
30   residue was chromatographed on a silica gel column using CHCl$_3$ as the eluent to give 51 mg (76%) of Va as yellow crystals: mp 213-215 °C; ^{1}H NMR (CDCl$_3$) δ 4.08, 4.10 (2 x 3 H, 2 s, 1- and 8-OMe), 4.47, 4.54 (2 X 2 h, 2 s, CH$_2$Br), 7.24-8.04 (10 H, m, H-2,4,5,6,7). Anal. (C$_{16}$H$_{11}$BrO$_4$) C, H, Br.

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EXAMPLE 11

3-(Dibrom methyl)-1(and 8)-hydroxy-8(and 1)-methoxy-9,10-anthraquinone (VIIa) and 6-(Dibromomethyl)-1(and 8)-hydroxy-3,8(and 1,3)-dimethoxy-9,10-anthraquinone (VIIb). In a similar manner, from IVa (200 mg, 0.453 mmol) and IVb (100 mg, 0.213 mmol), VIIa (82 mg, 42.5%) and VIIb (82 mg., 84.4%), respectively, were prepared (see Table III).

EXAMPLE 12

3-(Bromomethyl)-1,8-dihydroxy-9,10-anthraquinone (VIa). A mixture of III a (1.05 g, 2.9 mmol), HOAc (50 mL), and 30% HBr/HOAc (5 mL) was heated at 100 °C for 5 h. After the mixture was cooled, VIa was collected by filtration, washed with HOAc and H_2O, and then air-dried to give 879 mg (91%) of the product: mp 220-222 °C; ^1H NMR (CDCl$_3$) δ 4.47 (2 H, s, CH$_2$Br), 7.22-7.90 (5 H, m, H-2,4,5,6,7), 12.01, 12.04 (2 x H, 2 s, 1-OH, 8-OH). Anal. (C$_{15}$H$_9$BrO$_4$) C, H, Br.

EXAMPLE 13

3-(Dibromomethyl)-1,8-dihydroxy-9,10-anthraquinone (VIIIa) and 6-(Dibromomethyl)-3-methoxy-1,8-dihydroxy-9,10-anthraquinone (VIIIb). In a similar manner, IVa (237 mg, 0.54 mmol) and IVb (472 mg, 1 mmol) were converted into VIIIa (166 mg, 75%) and VIIIb (408 mg, 95%), respectively (Table III).

EXAMPLE 14

3-[(Diethylamino)methyl]-1,8-dimethoxy-9,10-anthraquinone (IXa-1). To a solution of IIIa (200 mg, 0.55 mmol) in DMF (10.0 mL) was added Et$_2$NH (5.0 mL), and the mixture was stirred at room temperature for 3 days. The mixture was partitioned between EtOAc (20 mL) and H_2O (20 mL). The aqueous layer was washed with EtOAc (20 mL). The combined EtOAc solutions were washed with H_2O (2 x 20 mL) and saturated NaCl (2 x 20 mL), dried (Na$_2$SO$_4$), and concentrated, and the residue was chromatographed on a silica gel column first with CHCl$_3$, which eluted the 3-(hydroxymethyl)

derivative (33 mg), followed by CHCl$_3$ containing 3% M OH.
The IXa (140 mg) that eluted from the column was converted
to the crystalline HCl salt (142 mg, 63%), mp 154-158 °C.

In a similar manner but by using the corresponding amines,
IXa-2, -5, and -6, were prepared (Table III). Also by use
of the same procedure starting from IIIb and the
corresponding amines, IXb-1, -2, -5, and -7 were synthesized
(Table III).

EXAMPLE 15

3-[(Diethylamino)methyl]-(or 8)-hydroxy-8(or 1)-methoxy-
9,10-anthraquinone (Xa-1). The HCl salt monohydrate of IXa
(100 mg, 0.25 mmol) was dissolved in a mixture of HOAc (5
mL) and 30% HBr/HOAc (0.4 mL), and the solution was stirred
at room temperature for 24 h. After concentration in vacuo,
the residue was partitioned between saturated NaHCO$_3$ (10 mL)
and CHCl$_3$ (10 mL). The CHCl$_3$ layer was dried (Na$_2$SO$_4$) and
concentrated and the residue chromatographed on a silica gel
column using CHCl$_3$-MeOH (30:1 v/v) to give Xa-1 as a glass,
which was dissolved in 1 N HBr (1 mL). Upon dilution of the
solution with EtOH (5 mL), the monohydrobromide of Xa-1 (82
mg) precipitated as yellow microcrystals, mp 225-227 °C dec.

In a similar manner, Xa-2-4 and Xb 1-4 were prepared (Table
III).

EXAMPLE 16

3-[[N,N-Bis(2-hydroxyethyl)amino]methyl]-1,8-dihydroxy-9,10-
anthraquinone (XIa-2). A mixture of VIa (456 mg, 0.73 mmol)
and bis(2-hydroxyethyl)amine (600 mg, 5.50 mmol) in DMF (20
mL) was stirred for 2 h and then partitioned between CHCl$_3$
(100 mL) and H$_2$O, (100 mL). The organic layer was
separated, washed H$_2$O, 3 X 50 Ml), dried (Na$_2$SO$_4$), and
concentrated in vacuo. The residue was chromatographed on
a silica gel column using CHCl$_3$-MeOH (15:1 v/v) as the
eluent. The major fraction was concentrated and the residue

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dissolved in 1 N HCl. After concentration of the solution
in vacuo, the residu was triturated well with MeOH (5 mL),
and the crystalline HCl salt of XIa-2 (496 mg, 91%) was
collected by filtration, mp 204-207 °C. Anal. ($C_{19}H_{19}NO_6HCl$)
C, H, N.

5

In a similar manner but by use of the corresponding amines,
XIa-1, -2, -5, -6, 8-11, and -13-15 were prepared. Also,
from VIb, by the same procedure, XIb-1, -2, -5, and -6 were
obtained (Table III).

EXAMPLE 17

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3-[[N,N-Bis(2-chloroethyl)amino]methyl]-1,8-dihydroxy-9,10-
anthraquinone (XIa-3). To a solution of XIa-2-HCl (1.05 g,
2.57 mmol) in dry DMF (40 mL) was added $SOCl_2$ (1.0 mL), and
the solution was stirred at room temperature for 1.5 h. The
solution was concentrated in vacuo (bath temperature <40°C),

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and the residue was cooled in an ice bath. Cold MeOH (10
mL) was added to destroy DMF-HCl complex, and the mixture
was concentrated in vacuo. Upon trituration of the residue
with cold meOH (10 mL), XIa-3 (977 mg, 85.1 %) was obtained,
mp 211-214 °C. Anal ($C_{19}H_{17}Cl_2NO_4HCl$) C, H, N.

20

In a similar manner but by use of the corresponding amino
alcohols, IXa-3 and -7, IXb-3 and -7, Xa-3, Xb-3, XIa-7, and
XIb-3 and -7 were synthesized (Table III).

25

Spectral Studies. Absorption spectra were measured with an
IBM 9410 UV-visible spectrometer interfaced to an HP 9826
computer. Small volumes of the stock drug solutions (2
mg/mL in Me_2SO) were added to 2 mL of buffer (0.01 or 0.1 M
NaCl, 5 mM Hepes, pH 7) to obtain a final drug concentration

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of 5-15 μM, or to the solution of native or thermally
denatured DNA (0.2 and 0.1 mM, respectively) in the buffer.
After incubation at room temperature for 10 min, the spectra
were recorded in the 300-600 nm range (increment 1 nm) and

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corrected by subtracting the spectrum of the blank which was
measured before addition of the drug.

Biological Assays. Method A. For cell growth inhibition
studies, HL-60 cells 92.0 x 10^5/mL) were grown in RPMI 1640
media at 37°C in humidified 5% CO_2 for 72 h. Viable cells
5   were counted with the trypan blue exclusion method. The
fractional inhibitions at four or five concentrations of
compounds (in 0.2% DMSO) were analyzed with a median-effect
plot (Chou, T.-C.;=, et al., Adv. Enzyme Regul. 22:27
(1984)) by using a computer program. (Chou, J.; Chou, T.-C.
10 Dose-Effect Analyses with Microcomputer. Quantitation of
ID_{50}, LD_{50}, Synergism, Anagonism, Low-dose Risk, Receptor-
Binding and Enzyme Kinetics; IBM-PC Series, Elsevier-
Biosoft, Elsevier Scientific: Cambridge, U.K., (1986)) The
median-effect concentration (ID_{50}) was automatically
15 determined for the $\underline{x}$ intercept of the median-effect plot.
Cell growth in the absence of a compound and in the presence
of DMSO was used as a control. DMSO (0.2%) alone inhibited
cell growth $3.8 \pm 1.2\%$ during the 72-h incubation period.

20 **Method B.** For precursor incorporation studies, each compound
at four to six concentrations (0.2% DMSO) was preincubated
with HL-60 cells (2.5 X 10^6/Ml) for 15 min prior to the
addition of [^{3}H-methyl]TdR (1 μCi, 0.15 nmol/mL) and was
incubated for 30 min. The incubation conditions and the
25 procedures for isolating the DNA fractions were described
previously (Chou, T.-C, et al., J. Cancer Res. 43:3074
(1983)). The incorporation of radioactivity in DNA in the
absence of an analogue in the presence of DMSO was used as
a control. The control value for incorporation into DNA was
30 8500 $\pm$ 300 cpm/10^6 cells.

EXPERIMENT 3:

This experiment concerns Topoisomerase II (Topo II)-mediated
35 DNA cleavage activity induced by chrysophanol derivatives.

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3-(bis[2-chloroethyl]amino)methyl-1,8-dihydroxy-9,10-anthraquinone (SK-31662) and 3-(2-hydroxyethylamino)methyl-1,8-dihydroxy-9,10-anthraquinone (SK-31694) have antileukemic activity with IC_{50}s of 0.14 and 0.86 uM, respectively. SK-31662 and 31694 inhibit kDNA decatenation at 48 and 38 uM, respectively. The mapping of DNA-Topo II cleavage sites using Hind III-digested 3'-labeled DNA and nuclear extracts (NE) of HL-60 cells showed that at 10 uM these two agents induce protein-linked DNA breaks with a cleavage site pattern similar to m-AMSA. They also stimulate the formation of Topo II cleavable complex in the presence of 3'-labeled DNA and NE. The amounts of protein-linked DNA induced by VP-16 and SK-31694 are reduced during 0.5 min exposure to 65 degrees C whereas SK-31662-induced protein-linked DNA complex cannot be reversed up to 15 min. The data suggest that Topo II appears to be a major cytotoxic target for these compounds and DNA intercalator with alkylating groups interact with Topo II system in an irreversible manner with enhanced toxicity.

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What is claimed is:

1.   A compound having the structure:
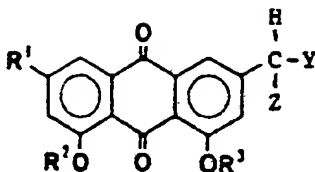
$$R^1 \text{—} \underset{R^2O \quad O \quad OR^3}{\overset{O}{\text{(anthraquinone ring system)}}} \text{—} \overset{H}{\underset{Z}{C}}\text{—}Y$$

wherein R^1 is hydrogen, a hydroxy group or a
methoxy group;
R^2 is hydrogen or a methyl group;
R^3 is hydrogen or a methyl group;
Y is a halogen, a secondary amino group or a
tertiary amino group and
Z is hydrogen or a halogen.

2.   A compound of claim 1, wherein the halogen is chlorine
or bromine.

3.   A compound of claim 1, wherein the secondary amino
group comprises a lower alkyl group or a substituted
lower alkyl group where each substituent is a halogen
or a hydroxy, benzoxy, or acetyloxy group.

4.   A compound of claim 1, wherein the tertiary amino group
comprises two lower alkyl groups or substituted lower
alkyl groups where each substituent is a halogen or a
hydroxy, benzoxy, or acetyloxy group and where the
lower alkyl groups or the substituted lower alkyl
groups are the same or different.

5.   A compound of claim 1 selected from the group
      consisting of:
      3-Bromomethyl-1,8-dihydroxy-9,10-anthraquinone;
      3-Dibromomethyl-1,8-dihydroxy-9,10-anthraquinone;
      3-Bromomethyl-1,8-dimethoxy-9,10-anthraquinone;
      3-Dibromomethyl-1,8-dimethoxy-9,10-anthraquinone;
      3-Bromomethyl-1,8-dihydroxy-6-methoxy-9,10-
            anthraquinone;
      3-Dibromomethyl-1,8-dimethoxy-6-methoxy-9,10-
            anthraquinone;
      3-Bromomethyl-1,6,8-trimethoxy-9,10-anthraquinone;and
      3-Dibromomethyl-1,6,8-trimethoxy-9,10-anthraquinone.

6.   A compound of claim 1 selected from the group
      consisting of:
      3-(bis[2-chloroethyl]amino)methyl-1,8-dihydroxy-9,10-
            anthraquinone; and
      3-(2-hydroxyethylamino)methyl-1,8-dihydroxy-9,10-
            anthraquinone.

7.   A compound of claim 1 selected from the group
      consisting of:
      3-(N,N-Diethylamino)methyl-1,8-dihydroxy-9,10-
            anthraquinone;
      3-(N-Ethylamino)methyl-1,8-dihydroxy-9,10-
            anthraquinone;
      3-[N,N-Bis(2-hydroxyethyl)amino]methyl-1,8-dihydroxy-
            9,10-anthraquinone;
      3-[N-(2-Hydroxyethyl)amino]methyl-1,8-dihydroxy-9,10-
            anthraquinone;
      3-(N,N-Diethylamino)methyl-1,8-dihydroxy-6-methoxy-
            9,10-anthraquinone;
      3-(N,Ethylamino)methyl-1,8-dihydroxy-6-methoxy-9,10-
            anthraquinone;
      3-[N,N-Bis(2-hydroxyethyl)amino]methyl-1,8-dihydroxy-
            6-methoxy-9,10-anthraquinone;

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3-[N-(2-hydroxyethyl)amino]methyl-1,8-dihydroxy-6-
     methoxy-9,10-anthraquinone;

3-(N,N-Diethylamino)methyl-1,8-dimethoxy-9,10-
     anthraquinone;

3-(N,Ethylamino)methyl-1,8-dimethoxy-9,10-
     anthraquinone;

5   3-[N,N-Bis(2-hydroxethyl)amino]methyl-1,8-dimethoxy-
     9,10-anthraquinone;

3-[N-(2-Hydroxethyl)amino]methyl-1,8-dimethoxy-9,10-
     anthraquinone;

3-(N,N-Diethylamino)methyl-1,6,8-trimethoxy-9,10-
10   anthraquinone;

3-(N-Ethylamino)methyl-1,6,8-trimethoxy-9,10-
     anthraquinone; and

3-[N,N-Bis(2-hydroxethyl)amino]methyl-1,6,8-trimethoxy-
     9,10-anthraquinone.

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8.   A compound of claim 1 selected from the group
     consisting of:
     3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-
         9,10-anthraquinone;
20   3-[N-(2-Chloroethyl)amino]methyl-1,8-dihydroxy-
         9,10-anthraquinone;
     3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dihydroxy-6
     methoxy-9,10-anthraquinone;
     3-[N-(2-Chloroethyl)amino]methyl-1,8-dihydroxy-6-
25   methoxy-9,10-anthraquinone;
     3-[N,N-Bis(2-chloroethyl)amino]methyl-1,8-dimethoxy-
         9,10-anthraquinone;
     3-[N-(2-Chloroethyl)amino]methyl-1,8-dimethoxy-
         9,10-anthraquinone;
30   3-[N,N-Bis(2-chloroethyl)amino]methyl-1,6,8-trimethoxy-
     9,10-anthraquinone; and
     3-[N-(2-Chloroethyl)amino]methyl-1,6,8-trimethoxy-
         9,10-anthraquinone.

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9. A pharmaceutical composition for the treatment of a malignancy in a subject comprising the compound of claim 1 or a pharmaceutically acceptable acid salt thereof and a pharmaceutically acceptable carrier, the amount of the compound or salt being 1 to 200 mg/kg of body weight of the subject.

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10. A method of treating a subject having a malignancy which comprises administering to the subject an effective amount of the compound of claim 1 to suppress growth of the malignancy.

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11. A method of claim 9, wherein the malignancy is leukemia.

12. A method of claim 9, wherein the malignancy is a tumor.

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1/6

Figure 1
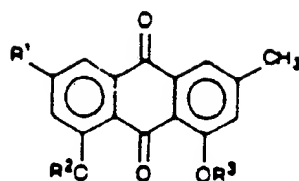
Ia R¹, R², R³ = H (chrysophenol)
Ib R¹ = OH, R², R³ = H (emodin)
IIa R¹ = H, R², R³ = Me
IIb R¹ = OMe, R², R³ = Me
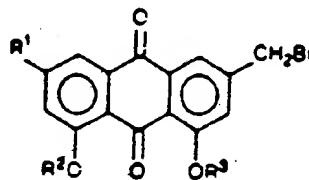
IIIa R¹ = H, R², R³ = Me
IIIb R¹ = OMe, R², R³ = Me
  Va R¹ = H, R² = H(Me), R³ = Me(H)
  Vb R¹ = OMe, R² = H(Me), R³ = Me(H)
 VIa R¹, R², R³ = H
 VIb R¹ = OMe, R², R³ = H
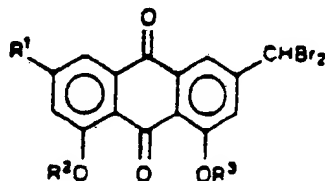
IVa R¹ = H, R², R³ = Me
IVb R¹ = OMe, R², R³ = Me
VIIa R¹ = H, R² = H(Me), R³ = Me(H)
VIIb R¹ = OMe, R² = H(Me), R³ = Me(H)
VIIIa R¹, R², R³ = H
VIIIb R¹ = OMe, R², R³ = H
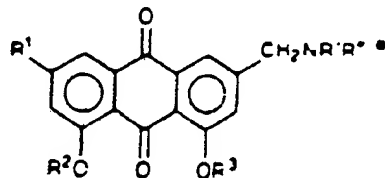
IXa R¹ = H, R², R³ = Me
IXb R¹ = OMe, R², R³ = Me
 Xa R¹ = H, R² = H(Me), R³ = Me(H)
 Xb R¹ = OMe, R² = H(Me), R³ = Me(H)
XIa R¹, R², R³ = H
XIb R¹ = OMe, R², R³ = H

*See Table III for NR'R''
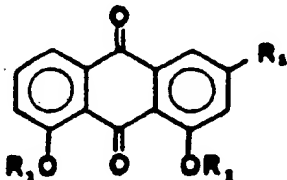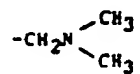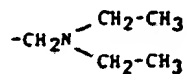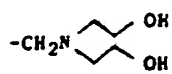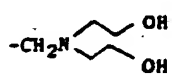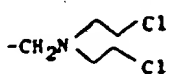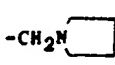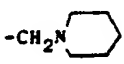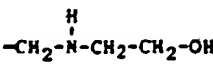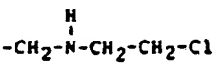
Figure 2

Potential Anticancer Agents
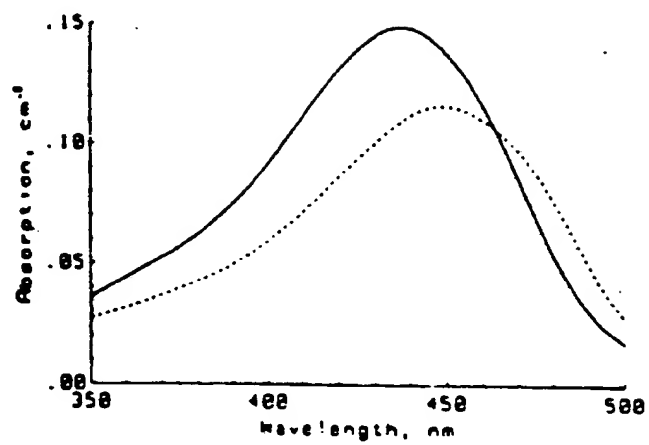
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Figure 3

Biological Effects of Chrysophanol Derivatives

| Compound	R1	R2	R3	IC$_{50}$ for cell growth (μM)
SK 31671	$-CH_2N(CH_3)_2$	H	H	2.8
SK 31661	$-CH_2N(CH_2-CH_3)_2$	H	H	1.8
SK 31660	$-CH_2N$ (OH, OH)	H	H	3.3
SK 31653	$-CH_2N$ (OH, OH)	H	CH$_3$	20.7
SK 31662	$-CH_2N$ (Cl, Cl)	H	H	0.14
SK 31669	$-CH_2N$ (azetidine ring)	H	H	1.14
SK 31665	$-CH_2N$ (piperidine ring)	H	H	2.8
SK 31694	$-CH_2-\overset{H}{N}-CH_2-CH_2-OH$	H	H	0.86
SK 31690	$-CH_2-\overset{H}{N}-CH_2-CH_2-Cl$	H	H	7.5
SK 31666	$-CH_2Br$	H	H	7.1

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Figure 4

TOPO II CLEAVABLE COMPLEX FORMATION
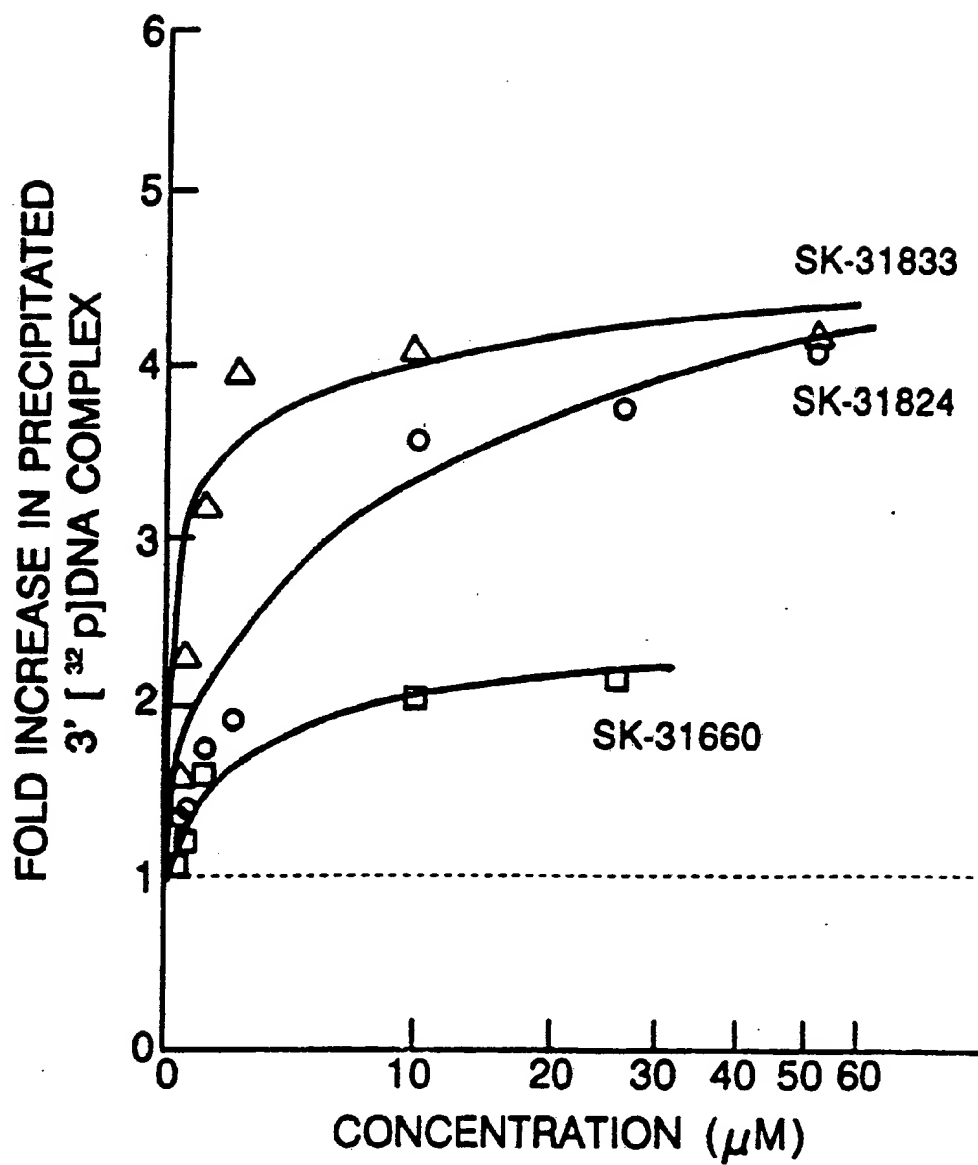
Figure 5

TOPO II CLEAVABLE COMPLEX FORMATION
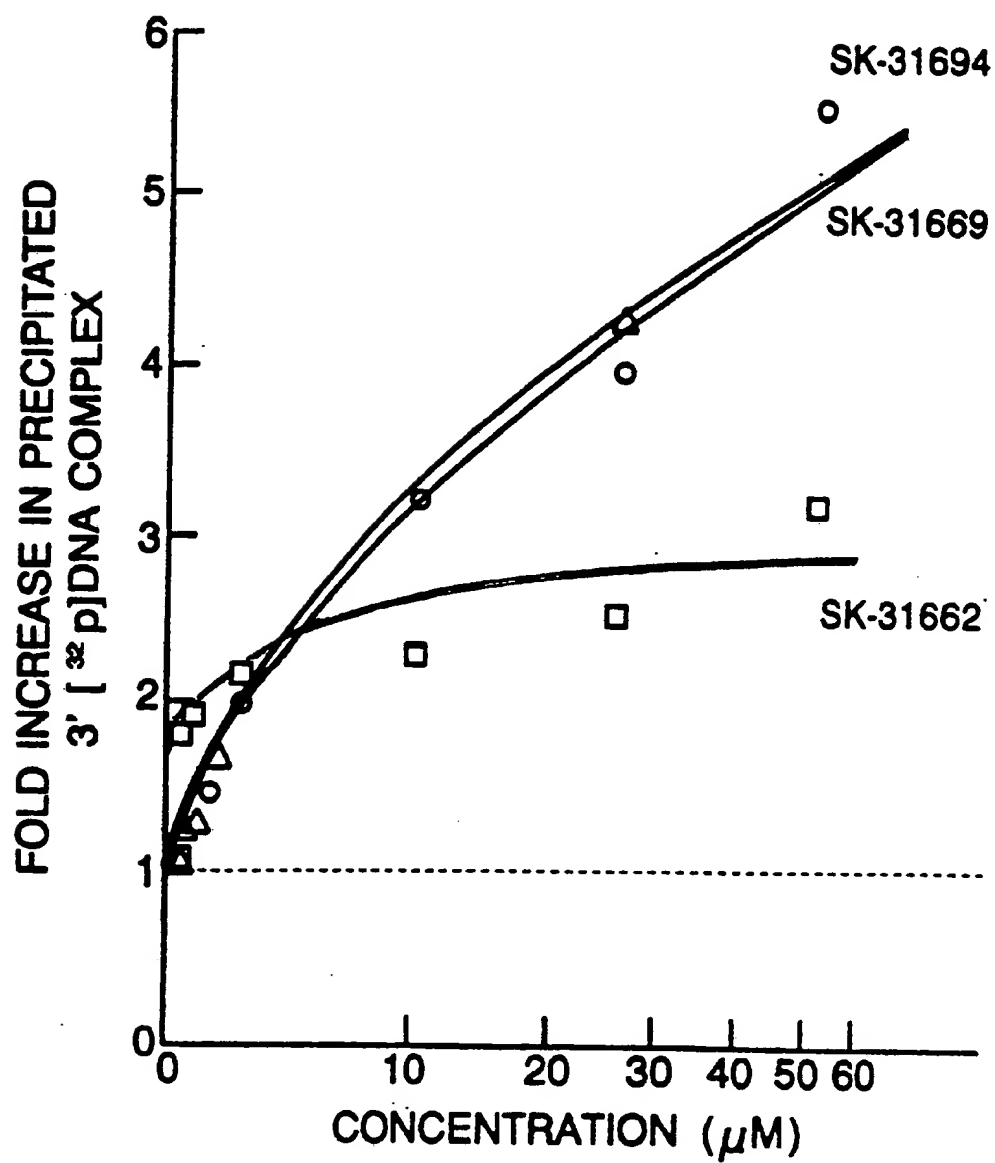
Figure 6

TOPO II CLEAVABLE COMPLEX FORMATION
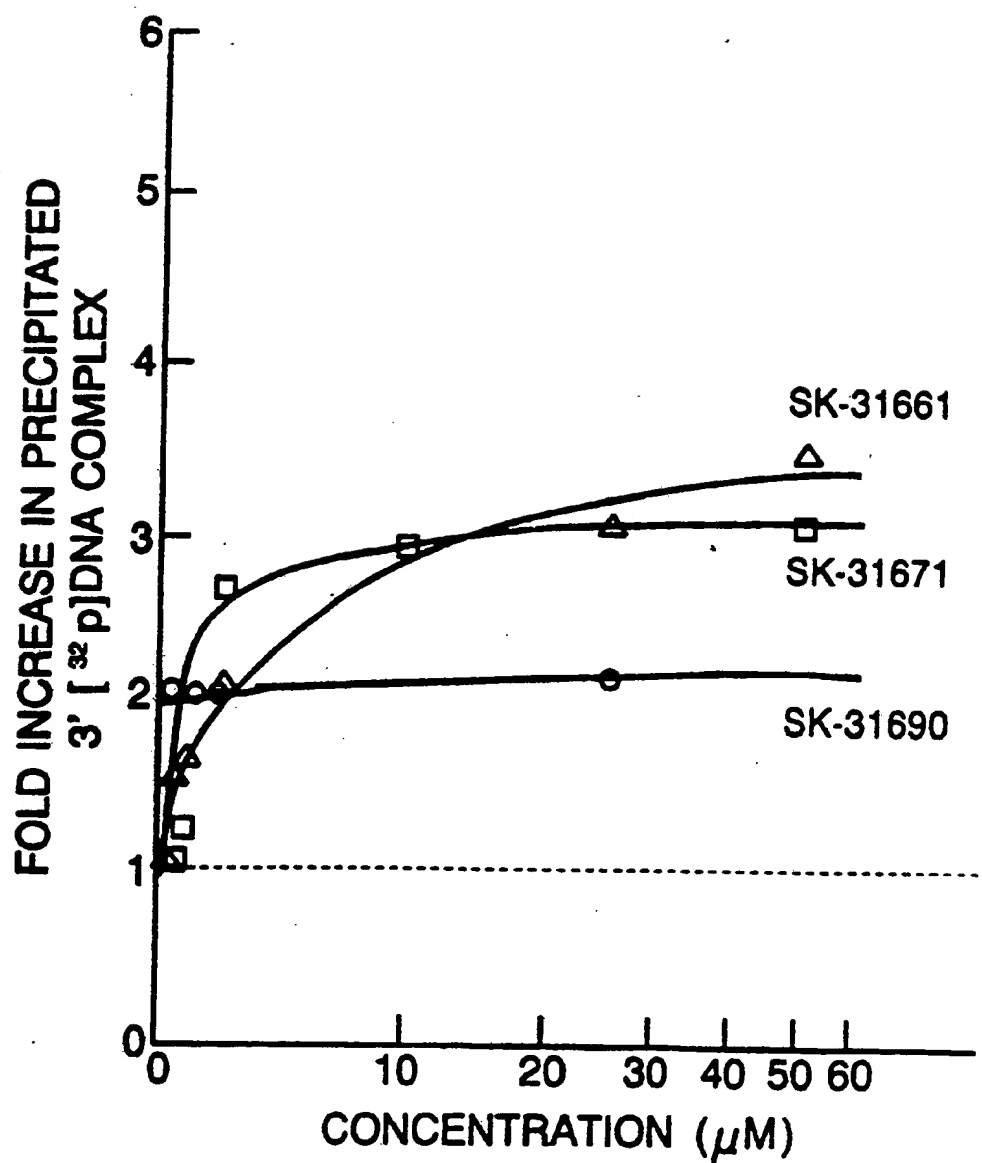
INTERNATIONAL SEARCH REPORT

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) 6

According to International Patent Classification (IPC) or to both National Classification and IPC

IPC(5): C07C 50/18, 34; A61K 31/03, 09, 12, 135
U.S. CL.: 552/261, 262, 266; 514/676, 680

II. FIELDS SEARCHED

Minimum Documentation Searched 7

Classification System	Classification Symbols
US, CL.	552/261, 262, 266; 514/676, 680

Documentation Searched other than Minimum Documentation
to the Extent that such Documents are Included in the Fields Searched 8

CAS Online: Graphic Structyre

III. DOCUMENTS CONSIDERED TO BE RELEVANT 9

Category *	Citation of Document, 11 with indication, where appropriate, of the relevant passages 12	Relevant to Claim No. 13
Y	US, A, 4,215,062 MITSCHER, 29 JULY 1980 See col. 4, lines 32-44.	1-12
Y	The Chemistry of Synthetic Dyes; Venkatoraman, "Natural Anthraquinones coloring Matters color Matters," Academic Press inc., New York, 1952 page 831.	1-12 1-12
Y	Anthracene and Anthraquine, Barnett, "The Halogen Compounds", D VAN NOSTRAND Company, New York, 1921, pages 174-175	1-12

* Special categories of cited documents: 10

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report
12 APRIL 1990	0 4 MAY 1990

International Searching Authority	Signature of Authorized Officer
ISA/US	RAYMOND COVINGTON

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V.☐ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE¹

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1.☐ Claim numbers _____ , because they relate to subject matter ¹² not required to be searched by this Authority, namely:

2.☐ Claim numbers _____ , because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out ¹³, specifically:

3.☐ Claim numbers _____ , because they are dependent claims not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI.☒ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING²

This International Searching Authority found multiple inventions in this international application as follows:

I.   Claims 1-9 drawn to Chrysophanol products.

II.   Claims 10-12 drawn to a method for treating maligancy.

1.☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.

2.☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:

3.☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

1 - 9

4.☐ As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

☐ The additional search fees were accompanied by applicant's protest.

☐ No protest accompanied the payment of additional search fees.

Form PCT/ISA/210 (supplemental sheet (2) (Rev. 11-87)